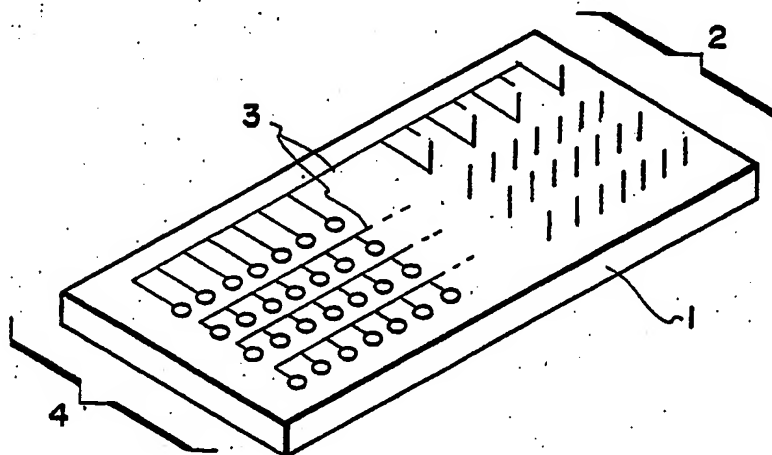




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(54) Title: ELECTRODE ARRAY AND METHOD OF MANUFACTURE



(57) Abstract

The electrode array is a device for making multiple electrical contacts with cellular tissue or organs. The electrode array includes a base (1), a two dimensional array of conducting protuberances (2) arising from the base and serving as electrodes, and conductors (3) embedded onto the base and connected to such protuberances for transmitting electrical signals to and/or from the protuberances. The protuberances may also include an insulating layer (15) which covers either the entire protuberance or which leaves the tips exposed for making focused electrical contact. Electrode arrays may be used singly or in combination with a second electrode array so as to form a sandwich around a target tissue. The sandwich electrode array (16, 17) may employ indexing cones for aligning the opposing electrode arrays and for limiting their vertical proximity. The conductors of the electrode array may be electronically connected or coupled to processing circuitry which amplifies and analyzes the signal received from the tissue and/or which generates signals which are sent to the target tissue and possibly coordinates the generated signals with signals which originate with the tissue.

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ELECTRODE ARRAY AND METHOD OF MANUFACTURE

RELATED APPLICATIONS

This application claims priority from U.S. Application No. 875,334, filed June 17, 1986, whose disclosure is, by reference, incorporated herein.

BACKGROUND

The invention relates to electrodes employed for electrically sensing or stimulating biological tissues. In particular, the invention relates to two dimensional electrode arrays and to methods for making and using such electrode arrays. The electrode array is particularly useful for making multiple electrical contacts at the cellular level, for electronically discriminating amongst individual cells or small groups of cells within a tissue or organ, and for directing electrical signals to or from such individual cells or small groups of cells within such tissue or organ, especially neural tissues and organs.

A nerve is a cordlike structure which is composed of numerous nerve fibers conveying impulses between a part of the central nervous system and some other region of the body. A nerve is made up of individual nerve fibers with their sheaths and supporting cells, small blood vessels, and a surrounding connective tissue sheath.

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Each nerve fiber is surrounded by a cellular sheath (neurilemma) from which it may or may not be separated by a laminated lipo-protein layer (myelin sheath). A group of such nerve fibers surrounded by a sheet of
5 connective tissue (perineurium) is called a fasciculus. The fasciculi are then bound together by a thick layer of connective tissue (epineurium) to form the nerve.

Neurologists have long sought an electrode device which could establish stable electrical contact with a
10 large number of individual nerve fibers within a nerve. Such a device would find wide medical application for sensing neurological impulses, facilitating the analysis and interpretation of such impulses, and delivering
15 electrical stimuli to target nerve fibers as a reaction to such analysis or as a result of external input. The ideal electrode device would be adapted to the anatomy of the nerve so that it could penetrate the nerve in a nondestructive fashion in order to form focused electrical contacts with a very large number of
20 individual nerve fibers.

Nerve cuff electrodes are employed in the neurological sciences for sensing nervous impulses and for electrically stimulating nerves. The nerve cuff electrode encircles the entire nerve and senses gross
25 nervous impulses arising from the nerve fibers within the nerve. The nerve cuff electrode may also be employed to electrically stimulate the nerve. Individual nerve fibers within a nerve may be functionally distinct from the other nerve fibers. The

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utility of the nerve cuff electrode is limited by its inability to specifically direct signals to or from selected nerve fibers within the nerve.

In order to make electrical contact with individual nerve fibers within a nerve, narrow gauge needle electrodes may be employed. When a narrow gauge needle is inserted into the nerve, there is a chance that it may make electrical contact with an individual nerve fiber or a small number of such fibers. If electrical contact is desired with each of several nerve fibers, then several needle electrodes must be employed. However, the technique of using multiple needle electrodes becomes progressively more and more difficult as the number of electrodes increases. Hence, there is a limit to the number of needle electrodes which can be usefully employed on a single nerve. Also, the electrical contact between a needle electrode and its corresponding nerve fiber can be disrupted by muscle motion and other forms of motion, since the end of the needle opposite the electrode extends outside the nerve and can be levered by relative motion of neighboring tissues. Therefore, long term implantation of needle electrodes with stable electrical contact with nerve fibers is not possible with prior art needle electrodes.

25 An electrode array having several electrodes integrated into one device is disclosed by Robert L. White. (Proceedings of the first International Conference on Electrical Stimulation of the Acoustic Nerve as a Treatment for Profound Sensorineural Deafness

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in Man, published by Velo-Bind, Inc. (1974), edited by Michael M. Merzenich, et al., chapter entitled "Integrated Circuits and Multiple Electrode Arrays," pp. 199-207, by Robert L. White) White's electrode array
5 employs a prong shaped base fabricated from a silicon wafer. The silicon base supports an array of electrodes which are deposited thereon toward the end of the prong. Each of the electrodes is small, flat, and circular, about 50 micrometers in diameter. Each electrode is
10 connected to a corresponding conductor which carries signals to and from the electrode. The conductor is electrically insulated from the tissue by a layer of silicon dioxide. In use, the prong is inserted tip first into neural tissue. Neural tissue is displaced by
15 the prong as it is inserted. Substantial damage to neural tissue can result from the insertion process due to the relatively large bulk of the prong. Since neural tissue slides tangentially past the electrodes during the insertion process, the flatness of the electrodes
20 helps to minimize the resultant disruption and destruction of neural tissue. However, once the device is inserted, the flatness of the electrodes limits the contact between the electrode and the neural tissue. Flat electrodes can make electrical contact only with
25 neural tissue which is directly adjacent to the surface of the prong.

A multiple electrode device with protruding electrodes is disclosed by Ko (IEEE Transactions on Biomedical Engineering, vol. BME-33 (3), pp 153-162

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(Feb. 1986), "Solid State Physical Transducers for Biomedical Research.") Ko discloses a device having three, sharp tipped, gold electrodes which protrude from the edge of a silicon base for making extracellular measurements. Like the White device, the silicon base of the Ko device has the shape of a prong and may cause significant tissue damage to the nerve during the insertion process. Also, if the device is implanted, its large bulk compromises the stability of the electrical contact between the electrodes and individual target cells. Additionally, the thinness of the prong can make it susceptible to shear damage with side loading. On the other hand, unlike the White device, the electrodes of the Ko device protrude horizontally from the edge of the silicon base so as to form the furthestmost extensions of the tip of the prong. The protrusion of the electrodes enhances the range of their contact. Unfortunately, the number of useful electrodes which may be incorporated into the Ko device is inherently limited. Since the electrodes are aligned in a row along the edge of the silicon base, it is not possible to array the electrodes into a configuration with more than one dimension. However, there are many bio-medical applications which can usefully employ a two dimensional array of protruding electrodes.

What was missing from the prior art and what was needed by practicing neurologists was an implantable electrode device which could electrically contact a large number of individual cells within an organ or

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tissu for sensing and controlling various bodily functions. The individual contacts should each be focused within a small region so that they involve single cells only. However, the range of the contacts should extend over a relatively large two or three dimensional region within the organ or tissue. The electrodes of the device should make positive contact with target cells and should be electrically stable over long periods of time, even with recurrent movement in adjacent tissues. On the other hand, the device should be able to penetrate the target organ without being intrusive so that tissue damage to the target organ is minimal. The device should have a small volume and a robust construction for practical medical applications.

15 SUMMARY

The electrode array is a device for establishing stable electrical contact with biological tissues. In the preferred embodiment, the electrode array has a configuration for making multiple extracellular contacts and for conducting electrical signals to or from each cell with which there is contact. However, the electrode array can also be employed for measuring the voltage potential of the surface of organs and tissues, e.g. for EKG or EEG.

25 The electrode array includes a base of semiconducting or nonconducting material, a two dimensional array of conducting protruberances which arise from such base and serve as electrodes, and conductors incorporated onto the base and connected to the protruberances for carrying

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electrical signals to and/or from such protruberances. The invention also includes various embodiments of the electrode array and methods for using and fabricating such electrode arrays.

5 In a preferred embodiment of the electrode array, the protruberances are coated with an insulating layer of dielectric material, except for their tips. This feature narrows and focuses the contact area of each protuberance to a relatively small region and

10 facilitates the ability of the protuberance to contact single cells or small groups of cells. The average number of extracellular contacts per protuberance may be adjusted to one by adapting shape and height of the protuberances and the exposed surface area of the tips.

15 In an alternative embodiment, the electrode array is capacitive. In this embodiment, the entire length of the protuberances, including the tip, is covered with an insulating dielectric. Hence each protuberance makes capacitive contact with cellular tissue.

20 In yet another embodiment which is particularly well adapted for establishing multiple electrical contacts with a large number of nerve fibers, a combination of two electrode arrays are employed to form a sandwich on either side of a nerve or target organ.

25 The two electrode arrays are situated on opposing sides of the nerve with the protuberances facing toward the center. The two electrode arrays are then brought closer together until they both contact the nerve and the protuberances penetrate into the nerve for making

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electrical contact with individual nerve fibers. At this point the electrode arrays are joined together by a holding means. The combination electrode array is then supported by the nerve to which it is clamped. Since
5 electrical contact is made on both sides of the nerve, the sandwich electrode array will make approximately twice the number of electrical contacts as compared to a single electrode array. Also, electrical contact between the electrode array and the nerve is enhanced by the
10 fact that the electrode array is supported by the nerve to which it is attached. Each of the electrode arrays within the sandwich may be either the conductive type or the capacitive type.

The invention also includes various biomedical
15 applications for the different embodiments of the electrode array. The electrode array may be either implanted or attached to skin. An electrode array may be employed for measuring the voltage potential of individual cells or of the surface area of an organ.
20 However, in the preferred application, the electrode array is surgically implanted for establishing long term electrical contact with multiple cellular elements of an internal organ or tissue. The implanted electrode array may either electrically stimulate individual cells
25 within the target organ or may sense nervous impulses within individual cells. Under some circumstances, the electrode array may both sense and stimulate electrical activity. Also, the electrical activity may be amplified and/or analyzed. And finally, the stimuli may

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be electronically correlated with the activity of the target cells. Because the two dimensional array greatly increases the number of protuberance which may be incorporated into a single device, the complexity and
5 redundancy of the protuberances is greatly enhanced. Consequently, it is possible to establish multiple electrical contacts with relatively complex biological systems.

The invention also includes various special
10 procedures employed for the fabrication and subsequent use of the electrode array. Since several electrode arrays may be fabricated on a single wafer, it is useful to employ indexing cones which mark out the various electrode arrays. The indexing cones can have a shape
15 which is similar to the protuberances, but are greater in height. After the electrode arrays have been deposited onto the wafer and the various subsequent steps have been completed, the indexing cones may be used as an index for guiding the sawing of the wafer
20 into separate base pieces. The indexing cones may also be employed with the sandwich electrode array for aligning the two electrode arrays with one another and for controlling and limiting the proximity of opposing electrode arrays so as to avoid damaging the sandwiched
25 nerve by exerting excessive pressure.

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BRIEF DESCRIPTION OF THE DRAWINGS

Fig. 1 is a perspective view of one embodiment of the electrode array illustrating a semiconductor base, an array sharp protuberances arising from the base, and
5 corresponding terminals. The array of sharp protuberances illustrate the concept of "bed of nails."

Fig. 2 is a a schematic diagram of the the electrode array of Fig. 1 illustrating the path of the individual conductors which electrically connect each protuberance
10 to a corresponding terminal or bonding pad.

Fig. 3 is an enlarged view of a fragment of the electrode array of Fig. 1, illustrating the pyramidal shape of the protuberances and indicating typical dimensions for the height of the protuberances and the
15 distance between adjacent protuberances.

Fig. 4 is a perspective view of a section of an alternative embodiment of the electrode array having conical protuberances illustrating a deposition mask attached to a metallic film atop the base for growing
20 the conical protuberances.

Fig. 5 is a further schematic diagram of the electrode array of Fig. 1 illustrating the layout of the protuberances, terminals, and conductor.

Fig. 6 is a sectional view of the electrode array of
25 Fig. 4 illustrating the relationship between the conical protuberances and the deposition mask.

Fig. 7 is sectional view of an alternative embodiment of the electrode array illustrating a protuberance having a dielectric coat covering the protuberance, exclusive

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f the tip.

Fig. 8 is a perspective view of two electrode arrays forming a sandwich on either side of a flattened nerve.

5 Fig. 9 is a sectional view of the two electrode arrays of Fig. 8 illustrating interdigitated protuberances penetrating a nerve from opposite sides and electrically contacting individual nerve cells.

Fig. 10 is a sectional view of Schwann cells
10 enveloping unmyelinated nerve fibers illustrating the conductive tip of a protuberance from an electrode array lying in close proximity to a nerve fiber.

Fig. 11 is a sectional view of two myelinated nerve fibers having nodes of Ranvier illustrating the
15 conductive tips of protuberances of differing heights from an electrode array lying in close proximity to said nodes.

Fig. 12 is a side plane view of an alternative embodiment of the electrode array having a monolithic
20 base structure, protuberances, and several electronic devices.

Fig. 13 is schematic diagram of the electrode array of Fig. 12 illustrating the interconnections for outputting the signal of the protuberances. Included
25 are a transmitter and receiver for transmitting signals between the protuberances and an external unit.

Fig. 14 is a sectional view of the fragment of two electrode arrays shown in Fig. 15 indicating the relative position of the opposing indexing cones.

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Fig. 15 is a plane view of indexing cones from two opposing electrode arrays illustrating the aligning and vertical positioning of the two electrode arrays by means of the indexing cones.

5

DETAILED DESCRIPTION OF THE INVENTION

The invention is an electrode array which is to be applied to body tissue to provide an effective electrical connection therewith, whether for sensing or stimulating purposes. The electrode array provides a
10 multiple possibility of successful electrical contact, and is intended to cause minimal damage to the body tissue or upset to the body system. The electrode array includes an array of conductive protuberances which serve as electrodes. The protuberances arise from a
15 base and are connected by electrical conductors to terminals on the base. The terminals and conductors may be employed to connect individual protuberances or groups of protuberances of the electrode array to other electrical circuits.

20

If the electrode array is to be used for sensing low voltage body signals, an amplifier would likely be the first electrical circuit connected to the protuberances and/or terminals. Then, of course, the signals (information) may go on to be handled by analog or
25 digital electronic methods and may involve transmission, multiplexing, filtering, data processing or other known electronic techniques. The particular use would

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determine the particular ther electrical circuits to be used.

If the electrode array is to be used for electrically stimulating a tissue, the terminals would be connected
5 to circuits which provide the output for the stimulation signals. The conductors would then carry these stimulation signals from the terminals to the corresponding protuberances.

Fig. 1 is a perspective view illustrating the concept
10 of a "bed of nails," showing the protuberances and terminals. It is drawn to illustrate the concept of a base (1) having protuberances (2) arising therefrom with conductors (3) leading from the protuberances (2) to terminals (4). The terminals illustrated in Fig. 1 are
15 bonding pads.

Fig. 2 is a more detailed view of Fig. 1 and illustrates the concept of connecting the an array of protuberances (2) to an array of terminals (4) by means of conductors (e.g. 5 & 6)

20 Fig. 3 is a view of an array of protuberances in the shape of pyramids, illustrating the dimensions which may be involved. The protuberances, or needles, may, of course, be taller and narrower. Spacing may vary, as may the size of the protuberances. Of course, such
25 protuberances may be conical or other elongated shapes.

Fig. 4 illustrates protuberances being grown through a mask onto a metallic film (9). The protuberances shown in Fig. 4 have the shape of cones or needles. Below the mask lies a sandwich which includes a silicon

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base (7), an insulation layer of silicon dioxide (8), and the metallic layer (9) upon which the protuberances are being grown. Above the metal layer (9) is a spacing layer (10). The spacing layer (10) may have a composition of silicon dioxide, photoresist, or other material. The spacing layer (10) is not required for all applications. Atop the spacing layer (10) is a the top mask or fine mesh screen (11). After the protuberances are completely grown, the mask is carefully removed, leaving the protuberances atop the metallic layer (9). The conductors are subsequently formed from the metallic layer (9).

Fig. 5 shows a schematic layout for an electrode array. An array of protuberances arise from a base (1) and are connected by electrical conductors (3) to bonding pads (4).

Fig. 6 is a cross-section of a deposition mask (11), showing the cones having been deposited through the holes of the mask. The cones (e.g. 12 & 13) are shown atop metallic layer (9). The underlying insulating layer (8) and base or substrate (7) are also shown.

Fig. 7 is an illustration of a needle protuberance (14) covered with an insulating layer of dielectric (15), e.g. silicon dioxide. The tips of the needle protuberances are left exposed and uncovered by dielectric (15). Below the protuberance (14) is metallic layer (9) upon which the conductors are formed. The underlying insulation layer of dielectric (8), e.g.

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silicon dioxide, is also shown. The underlying base is not shown.

Fig. 8 is an illustration of a combination of two electrode arrays (16 & 17) disposed on a single nerve (18) to form a sandwich electrode array or combination electrode array. The nerve is shown simply flattened although it may be further prepared to receive a sandwich electrode array by removal of a portion of its sheath and/or surrounding structures. The bonding pads or terminal portion of the electrode array may overhang from the nerve so as to clear the nerve in order to permit the bonding pads or terminals to be connected to external circuits. In one embodiment of the electrode array, the bonding pads or terminals are located on the edge of the base so as to facilitate the connection between the electrode array and external circuits.

Electrode arrays may be employed for measuring the voltage potential of the skin surface, e.g. for electrocardiograph and electroencephalograph measurements. In such applications, the electrode array may either penetrate the skin or may be applied more lightly. By penetrating the skin, a better connection is obtained without the use of conductive ointments. In addition, a capacitive coupling may be obtained by having the protuberances entirely covered with a passivating layer (dielectric) and applied to penetrate the skin. Thus, if the protuberances are electrically joined, the surface areas of the protuberances become one capacitive plate of substantial area and th

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dielectric lies between such plate and the other plate of the capacitor, viz. the surface of the skin or body tissue which is being measured.

Fig. 9 is a cross-section of a nerve and shows
5 interdigitated needle shaped protuberances as might occur from the arrangement shown in Fig. 8. The interdigitated needles (e.g. 19 & 20) are shown penetrating a nerve from opposite sides and contacting or coming into near proximity to the myelinated or
10 unmyelinated fibers (21 & 22). The needles are shown penetrating the perineurial sheath (43) and the extraperineurial tissue (44). Some of such tissue may be removed in preparation for the application of the electrode arrays. It is noted that the needles are
15 shown as exposed only at their tips or ends. Such structure is particularly useful in sensing, in order to limit the sensed electrical activity to a single fiber or a few fibers. A larger portion of the needle may be exposed in stimulating situations. In order to enhance
20 the likelihood of successfully sensing or stimulating a particular nerve fiber within a particular type of nerve, the dimensions, needle length, exposed tip length, amount of interdigitation, and needle spacing of the electrode array may be adapted to the anatomy of
25 such nerve.

Fig. 10 shows Schwann cell structures (23 & 46) disposed around "C" class nerve fibers, such as (25). A needle shaped protuberance (24) is shown in close proximity to nerve fiber (25).

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Fig. 11 shows two nerve fibers (26 & 27), their nodes of Ranvier (28 & 29), and needles (30, 31, & 32) penetrating into the nerve. Needles (30 & 32) are in proximity to said nodes and would more likely pick up 5 electrical signals than would needle (31).

Fig. 12 illustrates a monolithic base structure (33) in which several active electronic devices (34, 35, 36, 37, & 38) are created and on which are created the protuberances (2), for penetrating the body tissue.

10 Fig. 13 shows the interconnected electronic devices for switching the output of a sensory device. The transmitter and receiver (38) are shown, for transmitting the sensed information and receiving information for controlling the multiplexor (36) and the 15 selective logic (34) of the sensing needles, or protuberances. Logic control (37) provides control over the multiplexor (36) and selective logic (34). In this manner external control may be exercised in order to select particular needles which are in suitable contact, 20 or proximity, to desired nerve fibers. Amplifiers (35) provide increased signal strength. Integrated circuit technology may be used to provide the desired interconnections. Further, it may be appreciated that the transmitter and receiver (38) may be other than 25 radio frequency. They may transmit and receive utilizing infrared, magnetic induction, reflected impedance, acoustic waves, volumetric conduction or any other suitable well-known means for transmitting and receiving information. Such transmitter and receiver

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may be powered from inside or outside of the body. The entire implanted electrode array may be powered from outside the body by power transferred into the body through the receiver. In this manner, one or more
5 electrode arrays could be coordinated to operate together or in response to one another. An electrode array implanted in the brain could, without any wires (tetherless), communicate and control an electrode array attached to a muscle, a nerve or other body part. An
10 electrode array or several electrode arrays attached to the motor cortex of the brain could transmit, in tetherless fashion, many channels of information to receiving body parts, such as muscles, to which electrode arrays are attached.

15 Fig. 14 illustrates indexing cones or aligning means. Three indexing cones (39, 40, & 41) arise from a first base piece which a single crosshatched indexing cone (42) descending from a second opposing base piece. The indexing cones from the first and second base pieces
20 intermesh. The crosshatched cone (42) may register and align a mask, cover or other item which overlies the second base piece.

Fig. 15 shows a side view of the indexing cones of Fig. 14. and illustrates how such indexing cones
25 intermesh so as to index or align two devices. Two or more of such groups of indexing cones would be used in accomplishing the registration. It is not believed alignment was been achieved previously using such microstructures.

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In the preferred use of the invention, the electrode array is connected to a nerve. A nerve is generally of linear shape, but does not ordinarily lie in a straight line. Considering the needles of the array to be

5 longitudinally disposed along the direction of the nerve, one or more needles along such longitudinal direction may make contact with the same or different nerve fibers. The needles most likely to be useful are those which touch or are in close proximity to the

10 desired fibers. Laterally spaced needles may also be found to have made contact with the same nerve fiber. Other laterally spaced needles may connect to nearby nerve fibers which may have the same or different signals. Reinforcement of the sensing of signals can

15 thus be obtained. Similarly, reinforcement of stimulation signals can thus be provided. From the explanation provided above, it can be seen that sensing or stimulation of the same or different nerve fibers is possible.

20 The smallest class of nerve fibers are unmyelinated "C" fibers. Adjacent fibers of this class appear, from our own observation, to be spaced from approximately 1/2 micrometer to 5 micrometers apart, center to center. Larger nerve fibers, e.g. "A" and "B" fibers, which are

25 usually myelinated (surrounded by a sheath) appear to be spaced approximately 10 micrometers to 50 micrometers from adjacent fibers. In addition, a thickness of connective tissue encloses all of the component fibers in a nerve. In order to penetrate the nerve or in order

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to enter the fiber bundle sufficiently, but not too much, the needles would be approximately 1/2 micrometer high to on the order of 100 micrometers high. In selecting the correct needle height, consideration has to be given to the sheaths, Schwann cells, and other tissue to be penetrated in order to contact the nerve fiber. Similarly, for other tissues, the depth of penetration desired would determine the height of the needles. If the needles are fabricated with optimal materials and geometry within the above described dimensions, emphasizing a small tip radius, narrow taper, spacing and length appropriate to the tissue involved, the likelihood of making electrical contact with a minimum of tissue damage is high.

Depending on the capability of creating long needles, it is desired to have them as long and narrow as possible. Aspect ratios (height to base) of 10 to 1 are readily achievable. A needle which is 100 micrometers high might have a base of from 5 micrometers to 10 micrometers in diameter or greater.

It should be appreciated that the small size of the needles minimizes the likelihood that nerves, organs, tissue, or other body parts would be damaged by application of the electrode array and penetration by the needles.

The spacing of the needles, transversely across a nerve, would be from approximately 1/2 micron to on the order of 100 micrometers. "On the order of" means, in this context, and as used herein, within the range of

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1/10 of the dimension to 10 times the dimension.

Spacing of the needles along the length of a nerve might well be greater than the lateral spacing of the needles across the nerve. That is, the spacing distance between
5 needles along the length of a nerve can vary a great deal. Needles or groups of needles might well be longitudinally spaced 1000 micrometers, 2000 micrometers, etc., from one another, depending on the
10 desired density of electrical contact with the nerve.

The needles (electrodes) must, therefore, be spaced having in mind the specific application. The needles should be small and sharp enough to avoid damaging the nerve. Also the electrically conductive portion of each
15 needle should be small enough to contact only a single fiber and thereby obtain signals from only one fiber. Consequently, a preferred embodiment of the invention is to insulate the needles, except at or near their tips so that only a small electrically conductive portion of
20 each needle is exposed. In this way, each needle is less likely to electrically contact more than one fiber.

In addition, the needles must be high or long enough to assure sufficient penetration of the desired nerve so as to make electrical connection with the nerve fiber
25 inside the nerve. In order to reach the nerve fiber, the sheath and other connective tissues must be penetrated. However, "electrical connection" or "contact" with a nerve fiber or other body tissue may mean actual physical contact with the nerve fiber or tissue or it may mean being in sufficiently close

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location to sense the electrical signals therefrom or to stimulate the fiber or tissue as discussed previously in connection with Fig. 11. Further, as discussed previously, if the needles are entirely covered with a dielectric and utilize capacitive coupling, the needles do not actually make conductive contact with the body tissue.

If the longitudinal direction of the electrode array is slightly canted with respect to a nerve, electrical contact by some of the needles with some of the nerve fibers is greatly enhanced.

The spacing and needle length may vary on a given base. In order to reach down into a fissure in the brain, for example, it may be desirable to have longer needles on one portion of the electrode array and shorter needles on another portion. Also, spacing density on one portion of the electrode array may be greater or lesser than on another portion. There may be an abrupt change of needle length or density, or both, in one or more directions. Or there may be a graded or gradual changes in one or more directions.

It is to be understood that the array may be sized to fit the particular application and may be planar, multiplanar, curved, twisted, or other desired shape as required in the particular circumstances involved.

Ordinarily, the needles of the electrode array would be disposed on a rigid base. However, it is to be appreciated that the base may be flexible, or that the electrode array may be comprised of needles on a

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plurality of bases. In general, the needles in an array should be held in relatively fixed spacing with respect to each other. It is intended to cover by "relatively fixed" terminology, instances in which the base is

5 flexible, curved, stretchable, etc. Among the suitable bases are silicon, sapphire, or germanium. Numerous ceramics are also suitable for such biomedical use. Biomedical grade plastics may also be used such as the polyamides, polymethacrylate, acrylics, polycarbonates,

10 etc., to the extent that such plastics may be implantable or rendered implantable.

The needles may be arranged in random fashion or ordered in columns and/or rows or other ordered arrangements. The optimum embodiment from the

15 standpoint of orderly electrical connection is an ordered arrangement. One embodiment which may be desired is that in which each electrode (except, of course, those near the edges of the array) is surrounded by six other electrodes, all equidistantly spaced. The

20 needles are electrically connected to a terminal which may, likewise, be randomly located or located in columns and/or rows. The terminal may include bonding pads which provide an electrical connection between the needles and other electrical circuits. Connection

25 points need not be in the same arrangement as the needles. Thus, the needles may be located in columns, but not rows, and the terminals may be located in columns and rows.

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It should be understood that the electrode array, as described herein, provides a greater likelihood than the prior art of successfully contacting a desired nerve fiber or desired location in a part of the brain or other part of the body. Through testing and selection of appropriate terminals, needles which have successfully made a desired contact with a particular nerve fiber or target cell can be connected to output equipment for sensing purposes or input equipment for stimulating purposes.

It may be further understood that the electrical parameters which govern the successful application of the electrode array, employed either as a recording electrode or as a stimulating electrode, are the same as the parameters employed for prior art electrodes. For stimulating, the parameters include stimulus rate, wave form, analog or pulsatile type, and amplitude sufficient to depolarize nearby neurons without exceeding the minimum amplitude sufficient to cause electrolysis at the electrode surface. For sensing, the parameters involve the reduction of noise and amplification of signal. These various electrical parameters are discussed in the prior art literature and may be employed for use and operation the electrode arrays disclosed and described herein.

The needles may be constructed as "cones" and a method of construction may use techniques similar to those taught in United States Patents Nos. 3,755,704, 3,789,471, and 3,812,559, each naming Charles A. Spindt

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et al. as inventors. United States Patent No. 3,453,478, naming Kenneth R Soulders and Louis N. Heynick as inventors, also discloses background technology for constructing cones. Of course, it is not
5 essential that the needles be "cones" as described therein, but may be of pyramidal shape or shaped as any sharp protuberance. Further information on the fabrication technology involved, may be found in an article by C. A. Spindt and others, entitled "Physical
10 Properities of Thin-Film Field Emission Cathodes with Molybdenum Cones," Journal of Applied Physics, vol. 47 (12), December 1976. In those patents and the article, the intended use of the structure and method is to provide field emission cathodes and field ionizers.
15 Such needles, as disclosed by Spindt, contemplate electron-emitting structures as may be utilized in a vacuum tube. Also, he contemplates an electric field of megavolts per centimeter and current density of millions of amperes per square centimeter. For electron
20 emission, contemplated voltages are of the order of kilovolts and for field ionization, approximately ten fold higher. See Col. 2, 1.3 et seq., patent No. 3,812,559.

The device of the invention, on the other hand, as
25 either a sensor or a stimulator, is concerned with very low electrical currents and voltages. The needles of the electrode array of this invention would, ordinarily, not be connected in common, but each needle would provide its individual output, although it is to be

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und rstood that groups of needles could be connected
tog ther, t pr vide a common r reinf rced output of
either stimulation or sensing. Further, in a particular
situation, all needles of an array could be connected
5 together to provide a single stimulating output or a
single sensing output.

In one contemplated method of manufacture, a common
base (substrate) is used in order to mount the needles
and to achieve desired deposition. The base may have to
10 be modified to provide the desired isolation of the
individual needles or needle groupings. Such original
base, as modified, may provide the necessary electrical
conductors to convenient terminals of bonding pads for
connecting to other electrical circuits.

15 The various steps of manufacture of the electrical
conductors and terminals (bonding pads) may be
accomplished by known techniques of chemical or
electrical plating, etching, diffusing, sputtering,
evaporation or other suitable techniques. This may be
20 accomplished by using photolithographic or photgraphic
techniques, masks, photoresists, etchants, and
associated materials, known to those skilled in the
microcircuit art.

A suitable mask may be generated by a drawing,
25 followed by a photograph of the drawing, the making of a
negative or positive, covering a mask material with a
photoresist, exposing the photoresist through the
negative or positive, developing it and etching to
generate the mask. Fine mesh screens may be readily

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purchased or a mask may be created as described above, or by other known techniques.

In one embodiment, the steps of manufacture are as follows:

- 5 1. A non-conductive substrate, e.g. silicon having a silicon dioxide layer formed thereon, is used. A foil or film of conductive material is affixed thereon, possibly by sputtering, evaporation or other known integrated circuit manufacturing technologies;
- 10 2. Using a photoresist and a suitable mask, a pattern of electrical conductors and terminals (bonding pads) is laid out on the conductive material and all the rest of the material is etched or removed. It would be possible to commence with a non-conducting
- 15 substrate, and using known chemical deposition techniques, lay down a sensitizer in the form of the desired conductive pattern, which would allow subsequent chemical deposition of a conductive metal as the electrical conductors and terminals;
- 20 3. After cleansing the article, a glass passivation layer is laid down on the electrical conductors and terminals;
4. Again, a photoresist, a suitable mask, defining the needle sites, and an etchant are used in order to
- 25 locate the needle sites and to etch through the glass passivating layer, exposing each of the sites for growing a needle on an electrical conductor of the layer below;

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5. The same mask or a similar mask having holes therethrough, at the desired needle sites is disposed over the exposed needle sites in registration with such sites, and deposition of the needles is accomplished through such mask by metallic evaporation using, for example, electron beam or resistive element heating, in a high vacuum chamber. The metal deposits on the mask as well as within the hole on the needle site. The size of the hole becomes progressively smaller as metal is deposited atop the mask. The reduction of the size of the hole is precisely correlated with a reduction in the rate of metal deposition within the hole. The reduction of the size of the hole also reduces the target field within the hole upon which the metal is deposited. As a result, the protuberance formed within each hole assumes a tapered shape, e.g. conical, pyramidal, or needle shaped. The evaporating metal used to form the cones (needles) may be platinum, activated iridium, platinum iridium alloy, possibly, rhenium, or other suitable implantable electrode material. It is desired that the cones be made of a conductor which can deliver stimulus current, if stimulating, or sense very small voltages, if sensing, with little or no corrosion. If the mask is a fine mesh screen through which the needles are deposited, the precise size of the holes required for creating the needles may be obtained by placing the mask (covering the device) in a vacuum

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deposition system and rotating the device about an axis vertical to its surface and depositing, at a grazing incidence, more metal on the screen or mask layer. This can be used to decrease the starting size
5 of the holes to any diameter. Upon arriving at the desired diameter, the needles may be created by orthogonally plating through such narrowed holes as taught in United States Patent No. 3,812,550, referred to above;

10 6. The mask through which deposition is accomplished is carefully removed, leaving the needles exposed and providing the "bed of nails;"

7. A photoresist, a mask having the pattern of the test points and terminals, and an etchant are used to
15 remove the passivating layer over the test points and terminals in order that connection can be made to the array; and

8.a. If it is desired to make a capacitive electrode array, the protuberances must be coated with a
20 passivating or insulating layer. Aluminum oxide ($Al(2)O(3)$) is a preferred composition for the passivating layer and is widely described and employed in the prior art for this purpose; or

8.b. If it is desired to make a conductive electrode
25 array, the focus and specificity of the protuberances can be enhanced by covering the protuberances with a passivating or insulating layer, except for an area of 1-5 square micrometers at the tips. Hence, electrical contact is made only at the tip of the protuberances

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and the probability of contacting only one cell is enhanced. The protuberances are initially covered over their entire height with a passivating layer, e.g. aluminum oxide ($Al(2)O(3)$). The passivating layer is then removed from a small area of the tips by exposure to a controlled plasma etch; or

8.c. Alternatively, passivation may be achieved by fabricating the protuberances with self passivating compositions or with a combination of self passivating and non-passivating composition. For example, the first 9/10ths of the height of the protuberances may be fabricated with tantalum, a self passivating composition. The incomplete cone will have a flat top and will form a passivating layer upon exposure to the atmosphere. However, before the passivation layer is allowed for form, the cone is then completed by the deposition of a non-passivating metal, e.g. gold, iridium, platinum, etc. The last 1/10th of the cone will remain conductive.

20 The above process utilizes various of the manufacturing steps disclosed in the above mentioned article from the Journal of Applied Physics and in the above mentioned patents.

The manufacturing operation may commence with a thin film sandwich of metal on a dielectric (e.g. silicon dioxide on a base of silicon). The conductive and terminal pattern is formed out of the metal layer, by etching away excess metal. Then the needles are deposited through an appropriately patterned mask t

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coincide with the conductive patterns, as desired.

After the needles have been formed, the entire device could be covered with a glass passivating coat, except with needle tips and terminals if they are desired to be
5 left exposed. They could, of course, be exposed later.

In another method, a thin film sandwich is used, having a bottom layer of dielectric, a next layer of metal, then a dielectric and then metal on top of that. The top layer of metal becomes the mask for creating the
10 needles. The thickness of the bottom dielectric layer is determined by what rigidity and strength is necessary in order to hold on to and carry the electrode array. The second dielectric thickness is determined by the spacing desired between the top metal layer (which will
15 form a mask for the needle growing) and the middle metal layer upon which the needles will be grown. A very thin second dielectric layer may be created between the metal layer by the use of evaporated silicon dioxide. The under layer of metal will form the needle sites, the
20 electrical conductors, test points, if any and terminals, (bonding pads, in one embodiment). The top layer of metal is used as a mask for depositing the needle cones on the under layer of metal. This is accomplished by first making holes in the top layer of
25 metal, at intended needle sites, without penetrating the dielectric between the metal layers. This is done by a selective metal etchant (together with a photoresist and a mask) which does not attack the dielectric. Then, an etchant is used to remove the dielectric between the

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metal layers, at the needle sites. The needles are then "grown" by vacuum evaporation, sputter ring or other known techniques. After having formed the needles on the metal layer on the bottom dielectric layer, all of the second dielectric layer and top metal layer would be removed. The excess metal, not needed for electrical conductors, test points and terminals, of the exposed under layer metal could then be removed. In the alternative, the entire underlayer metal could be removed and new metal, making electrical conductors between the needles and terminals could be deposited. The entire electrode array could then be covered with a passivating material, such as silicon dioxide, silicon nitride, aluminum oxide (Al_2O_3) or other biocompatible dielectric, and then selectively etched at the terminals, if desired and at the needle points.

If the substrate is silicon or germanium or the like, the electrical conductors and, if desired, switches, multiplexors, amplifiers and other electronic circuits may be provided by doping selected portions of the substrate or by other commonly used techniques. Electrical conductors may be created on the surface of the semiconductor material, in it, or through it, to the opposite side from the protuberances.

In obtaining registration or indexing of masks, covers, or other items, which must be aligned with the array, one or more groups of three cones or needles could be grown in two or more places on the array and a registering cone or needle grown on the other item to be

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aligned. A needle on the overlying device fits into the space within the group on the other device, as previously described in connection with Figs 14 and 15. Of course, the overlaying device may have the groups of
5 needles and the base have the single registering needles. Further, both devices may have a group which fits into a group on the other device.

The materials used in the structure must be biocompatible and suitable for use in or in connection
10 with the living body. It is understood, of course, that certain materials which are not considered biocompatible could be rendered suitable by being treated or covered with a biocompatible material. Thus, glass passivation (covering with glass), oxidation of certain materials,
15 the coating or depositing of biocompatible materials (such as, but not limited to, silicone rubber, certain metals and ceramics or one of the many plastics which are used in the body) may be used to provide a final product which is biocompatible and may be implanted.

20 The electrode or needle material may be platinum, activated iridium, a platinum iridium alloy, a conductive polymer, carbon or other suitable electrically conductive material known by those skilled in the art as suitable for use in connection with the
25 body. In general, metals or other conductive substances which are inert and are least subject to corrosion are used. In the case of stimulating devices, conductive materials which can handle the necessary current densities are required.

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In view of the above discussion, it may be understood that the electrode array would be useful in stimulating a gland or a nerve t or in the gland to cause the gland to be active or more active. The electrode array may be
5 used to cause hormonal secretions.

Other uses of a stimulating electrode array or a plurality of electrode arrays would include stimulation of a group of muscles or successive stimulation of groups or portions of a group in order to achieve a
10 desired muscular coordination. Such electrode array may be applied directly to or in the muscle or it may be applied to or in selected nerves (or the central or peripheral nervous system) to provide signals to the muscle. Also, a number of such electrode array may be
15 applied at different locations and their stimulation or sensing coordinated to achieve desired results.

One stimulation application of the electrode array or a plurality of such electrode arrays is in excitation of the brain to provide a sensory response, e.g. vision.
20 The electrode array and its numerous needles may be disposed in the visuosensory and visuopsychic areas of the brain, which involve several kinds of cells. The electrode array may be disposed along the optic nerve or the paths where the optic nerve enters the cortex. The
25 array may be attached to the cortex with the needles penetrating the brain rather than the optic nerve.

The concept of the invention in one of its more important aspects provides for electrical access to the

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individual elements of a tissue in order to determine which element or elements and its associated needle or needles are useful for the intended purpose. One or more needle outputs may be found to be useful in the
5 particular application involved.

It should also be appreciated that, as taught hereinabove, the device may be untethered, through one or more means for transmitting information, receiving information or receiving power.

10 Although the invention has been described and illustrated in detail, it is to be clearly understood that the same is by way of illustration and example only and is not to be taken by way of limitation, the spirit and scope of this invention being limited only by
15 the terms of the appended claims.

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What is claimed is:

1. An electrode array for establishing electrical contact with tissue of a living body, the electrode array comprising:
 - a base,
 - 5 a two dimensional array of electrically conductive protuberances supported by said base, each of said protuberances having a tip and extending from said base for electrically contacting the tissue,
 - a conductor incorporated onto said base and
 - 10 electrically connected to said protuberances for conducting electric signals to or from said protuberances, and
 - a dielectric coat covering and electrically insulating said protuberances, exclusive of the tips.
2. An electrode array as described in claim 1 wherein:
 - said protuberances perpendicularly extend from
 - said base with a range of heights from approximately 0.5
 - 5 micrometers to on the order of 100 micrometers.
3. An electrode array as described in claim 1 wherein:
 - said protuberances are adjacently spaced with a
 - range from approximately 0.5 micrometers to on the order
 - 5 of 100 micrometers upon said base.

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4. An electrode array as described in claim 1 for establishing an array of electrical contacts with tissue, wherein:

said conductor including a plurality of conductors
5 electrically connected to and corresponding to at least one of said protuberances for forming a two dimensional array,

whereby each tip is capable of establishing an electrical contact with the tissue and the two
10 dimensional array of said protuberances is capable of establishing an array of electrical contacts with the tissue.

5. An electrode array as described in claim 4 further comprising:

a plurality of means for generating a signal,
each of said generating means being electronically
5 coupled to and corresponding to one of said conductors and each of said conductors coupled to and corresponding to one of said generating means.

6. An electrode array as described in claim 4 further comprising:

means for analyzing a plurality of signals,
said analyzing means being electronically
5 coupled to and corresponding to each of said conductors.

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7. An electrode array as described in claim 4

wherein:

said protuberances perpendicularly extending from
said base with a range of heights from approximately 0.5
5 micrometers to on the order of 100 micrometers, and

said protuberances are adjacently spaced with a
range from approximately 0.5 micrometers to on the order
of 100 micrometers upon said base.

8. An electrode array as described in claim 4 wherein
the electrode array electrically contacts individual
cellular components of the tissue of the living body and
wherein:

5 each tip of said protuberances being sufficiently
small and sharp to be capable of making electrical
contact with a single cellular component within the
tissue.

9. An electrode array as described in claim 4
wherein:

said dielectric coat having a biocompatible
composition for rendering the electrode array
5 implantable into the living body.

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10. A capacitor electrode array for capacitively coupling with the tissue of a living body, the capacitor electrode array comprising:

a base,

5 a two dimensional array of electrically conductive protuberances supported by said base, each of said protuberances having a tip,

a dielectric coat covering said protuberances for capacitively coupling each of said protuberances with
10 the tissue, and

a conductor incorporated onto to said base and electrically connected to said protuberances, said conductor for conducting electric signals to or from said protuberances.

11. A capacitor electrode array as described in claim 10 for multiply capacitively coupling with tissue, wherein:

said conductor including a plurality of conductors electrically connected to and corresponding to at least
5 one of said protuberances,

whereby each tip is capable of capacitively coupling with the tissue and the two dimensional array of said protuberances is capable of multiply capacitively coupling with the tissue.

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12. A capacitor electrode array as described in claim 10 wherein:

said protuberances perpendicularly extending from said base with a range of heights from approximately 0.5 micrometers to on the order of 100 micrometers.

13. A capacitor electrode array as described in claim 10 wherein:

said protuberances are adjacently spaced with a range from approximately 0.5 micrometers to on the order of 100 micrometers upon said base.

14. A capacitor electrode array as described in claim 10 wherein:

said protuberances perpendicularly extending from said base with a range of heights from approximately 0.5 micrometers to on the order of 100 micrometers and

said protuberances are adjacently spaced with a range from approximately 0.5 micrometers to on the order of 100 micrometers upon said base.

15. A capacitor electrode array as described in claim 10 further comprising:

means for generating multiple electric signals,
said generating means being electronically coupled to said conductors.

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16. A capacitor electrode array as described in claim
10 further comprising:

means for analyzing multiple electrical signals,
said analyzing means being electronically coupled
5 to said conductors.

17. A combination of two electrode arrays for
establishing electrical contact with a nerve, the
combination comprising:

a first electrode array including a first base, a
5 first array of electrically conductive protuberances,
and a first set of conductors connected to the first
array of protuberances,

a second electrode array including second base, a
second array of electrically conductive protuberances,
10 and a second set of conductors connected to the second
array of protuberances, and

means for holding the first electrode array
opposite the second electrode array with the nerve
sandwiched between the first and second electrode arrays
15 and with the first and second array of protuberances
penetrating and contacting the nerve.

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18. A combination of two electrode arrays for capacitively coupling with a nerve, the combination comprising:

a first capacitor electrode array including a
5 first base, a first array of electrically capacitive protuberances having a dielectric coat, and a first set of conductors connected to the first array of protuberances,

a second capacitor electrode array including
10 second base, a second array of electrically capacitive protuberances having a dielectric coat, and a second set of conductors connected to the second array of protuberances, and

15 means for holding the first capacitor electrode array opposite the second capacitor electrode array with the nerve sandwiched between the first and second capacitor electrode arrays and with the first and second array of capacitive protuberances penetrating and
20 capacitively coupling with the nerve.

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19. A method for sensing the individual electrical activity of each individual cell of a plurality of cells belonging to an organ or tissue, the method comprising the following steps:

5 step (a): contacting the surface of the organ or tissue with an electrode array having a base, a two dimensional array of electrically conductive protuberances having tips, a set of conductors connected to the array of protuberances, and a dielectric coat
10 covering the protuberances, exclusive of the tips,

in said step (a), the tips making electrical contact with the plurality of individual cells within the organ or tissue, and.

15 step (b): sensing the individual electrical activity of each of the individual cells of the plurality of cells belonging to the organ or tissue by means of the tips of the electrode array.

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20. A method for electrically stimulating each individual cell of a plurality of cells belonging to an organ or tissue, the method comprising the following steps:

5 step (a): contacting the surface of the organ or tissue with an electrode array having a base, a two dimensional array of electrically conductive protuberances having tips, a set of conductors connected to the array of protuberances, and a dielectric coat
10 covering the protuberances, exclusive of the tips,

in said step (a), the tips making electrical contact with the plurality of individual cells within the organ or tissue, and

step (b): electrically stimulating one or more of
15 the individual cells of the plurality of cells belonging to the organ or tissue which are in electrical contact with one or more of the tips of the electrode array.

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21. A method for capacitively sensing the individual electrical activity of each individual cell of a plurality of cells belonging to an organ or tissue, the method comprising the following steps:

- 5 step (a): contacting the surface of the organ or tissue with an electrode array having a base, a two dimensional array of electrically conductive protuberances having tips, a set of conductors connected to the array of protuberances, and a dielectric coat
10 covering and imparting a capacitance to the protuberances,

 in said step (a), the tips capacitively coupling with the plurality of individual cells within the organ or tissue, and

- 15 step (b): capacitively sensing the individual electrical activity of each of the individual cells of the plurality of cells belonging to the organ or tissue by means of the protuberances of the electrode array.

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22. A method for capacitively stimulating each individual cell of a plurality of cells belonging to an organ or tissue, the method comprising the following steps:

5 step (a): contacting the surface of the organ or tissue with an electrode array having a base, a two dimensional array of electrically conductive protuberances having tips, a set of conductors connected to the array of protuberances, and a dielectric coat
10 covering and imparting a capacitance to the protuberances,

 in said step (a), the tips making capacitive electrical contact with the plurality of individual cells within the organ or tissue, and

15 step (b): capacitively electrically stimulating one or more of the individual cells of the plurality of cells belonging to the organ or tissue which are in capacitive electrical contact with one or more of the tips of the electrode array.

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23. A method for sensing the individual electrical activity of each individual cell of a plurality of cells belonging to an organ or tissue, the method comprising the following steps:

5 step (a): contacting the surface of the organ or tissue with a combination of two electrode arrays including a first electrode array, a second electrode array, and a holding means,

 the first electrode array including a first base,
10 a first array of electrically conductive protuberances, and a first set of conductors connected to the first array of protuberances,

 the second electrode array including second base,
a second array of electrically conductive protuberances;
15 and a second set of conductors connected to the second array of protuberances, and

 the holding means for holding the first electrode array opposite the second electrode array with the organ or tissue sandwiched between the first and second
20 electrode arrays and with the first and second array of protuberances penetrating and contacting the plurality of individual cells within the organ or tissue, and

 step (b): sensing the individual electrical activity of each of the individual cells of the
25 plurality of cells belonging to the organ or tissue by means of the combination of two electrode arrays.

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24. A method for electrically stimulating each individual cell of a plurality of cells belonging to an organ or tissue, the method comprising the following steps:

- 5 step (a): contacting the surface of the organ or tissue with combination of two electrode arrays including a first electrode array, a second electrode array, and a holding means,
- the first electrode array including a first base, 10 a first array of electrically protuberances, and a first set of conductors connected to the first array of protuberances,
- the second electrode array including second base, a second array of electrically protuberances, and a 15 second set of conductors connected to the second array of protuberances, and
- the holding means for holding the first electrode array opposite the second electrode array with the organ or tissue sandwiched between the first and second 20 electrode arrays and with the first and second array of protuberances penetrating and making electrical contact with the plurality of individual cells within the organ or tissue, and
- 25 step (b): electrically stimulating one or more of the individual cells of the plurality of cells belonging to the organ or tissue which are in electrical contact with the combination of two electrode arrays.

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25. A method for capacitively sensing the individual electrical activity of each individual cell of a plurality of cells belonging to an organ or tissue, the method comprising the following steps:

- 5 step (a): contacting the surface of the organ or tissue with a combination of two capacitive electrode arrays including a first capacitive electrode array, a second capacitive electrode array, and a holding means, the first capacitive electrode array including a
- 10 first base, a first array of electrically capacitive protuberances, and a first set of conductors connected to the first array of capacitive protuberances,
- the second capacitive electrode array including second base, a second array of electrically capacitive
- 15 protuberances, and a second set of conductors connected to the second array of capacitive protuberances, and
- the holding means for holding the first capacitive electrode array opposite the second capacitive electrode array with the organ or tissue sandwiched between the
- 20 first and second capacitive electrode arrays and with the first and second array of capacitive protuberances penetrating and contacting the plurality of individual cells within the organ or tissue, and

- step (b): sensing the individual electrical
- 25 activity of each of the individual cells of the plurality of cells belonging to the organ or tissue by means of the combination of two capacitive electrode arrays.

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26. A method for capacitively stimulating each individual cell of a plurality of cells belonging to an organ or tissue, the method comprising the following steps:

- 5 step (a): contacting the surface of the organ or tissue with combination of two capacitive electrode arrays including a first capacitive electrode array, a second capacitive electrode array, and a holding means, the first capacitive electrode array including a
- 10 first base, a first array of electrically capacitive protuberances, and a first set of conductors connected to the first array of capacitive protuberances,
- the second electrode array including second base, a second array of electrically capacitive protuberances,
- 15 and a second set of conductors connected to the second array of capacitive protuberances, and
- the holding means for holding the first capacitive electrode array opposite the second capacitive electrode array with the organ or tissue sandwiched between the
- 20 first and second capacitive electrode arrays and with the first and second array of capacitive protuberances penetrating and making capacitive contact with the plurality of individual cells within the organ or tissue, and
- 25 step (b): capacitively stimulating one or more of the individual cells of the plurality of cells belonging to the organ or tissue which are in capacitive contact with the combination of two capacitive electrode arrays.

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27. A method for making an electrode array comprising the following steps:

step (a): obtaining a base with a non-conductive surface, then

5 step (b): depositing a layer of electrically conductive material atop the non-conductive surface, and

step (c): defining sites upon the layer of electrically conductive material for an array of protuberances, and then

10 step (d): depositing electrically conductive protuberances upon the sites for the array, the protuberances having tips, and

step (e): forming conductors for electrically connecting to the protuberances by partially removing
15 the layer of electrically conductive material, and then

step (f): depositing a dielectric coat upon the protuberances, exclusive of the tips.

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28. A method for making a capacitive electrode array comprising the following steps:

step (a): obtaining a base with a non-conductive surface, then

5 step (b): depositing a layer of electrically conductive material atop the non-conductive surface, and

step (c): defining sites upon the layer of electrically conductive material for an array of protuberances, and then

10 step (d): depositing electrically conductive protuberances upon the sites for the array of protuberances, the protuberances having tips, and

step (e): forming conductors for electrically connecting to the protuberances by partially removing
15 the layer of electrically conductive material, and then

step (f): depositing a dielectric coat upon the protuberances.

29. An indexible wafer comprising:

a wafer and

an indexing cone arising from said wafer.

30. An indexible electrode array comprising:

a base,

one or more protuberances arising from said base,

and

5 one ore more indexing cone arising from said base, said indexing cone rising to a greater height from said base than said protuberances.

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31. A pair of indexible electrode arrays comprising:
two bases,
one or more protuberances arising from each of
said bases, and
5 one or more indexing cone arising from each of
said bases,
said indexing cones rising to a greater height
from their corresponding bases than said protuberances,
said indexing cones being positioned on each of
10 said bases for aligning and spacing the two electrode
arrays with respect to one another.

32. A method of aligning a wafer comprising:
aligning the wafer with an indexing cone.

33. A method of aligning and spacing two electrode
arrays with respect to one another, each electrode
array having one or more indexing cones, the method
comprising:
5 aligning the electrode arrays with the indexing
cones.

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34. An electrode array comprising:

a base,

multiple protuberances arising from said base, said
protuberances having a self passivating composition
5 proximate to said base and a non-passivating composition
distal to said base, and

conductors embedded onto said base and connected to
said protuberances.

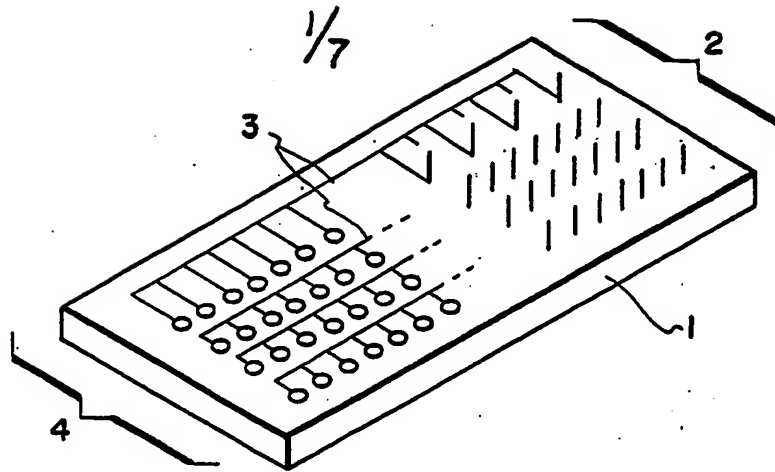


Fig. 1.

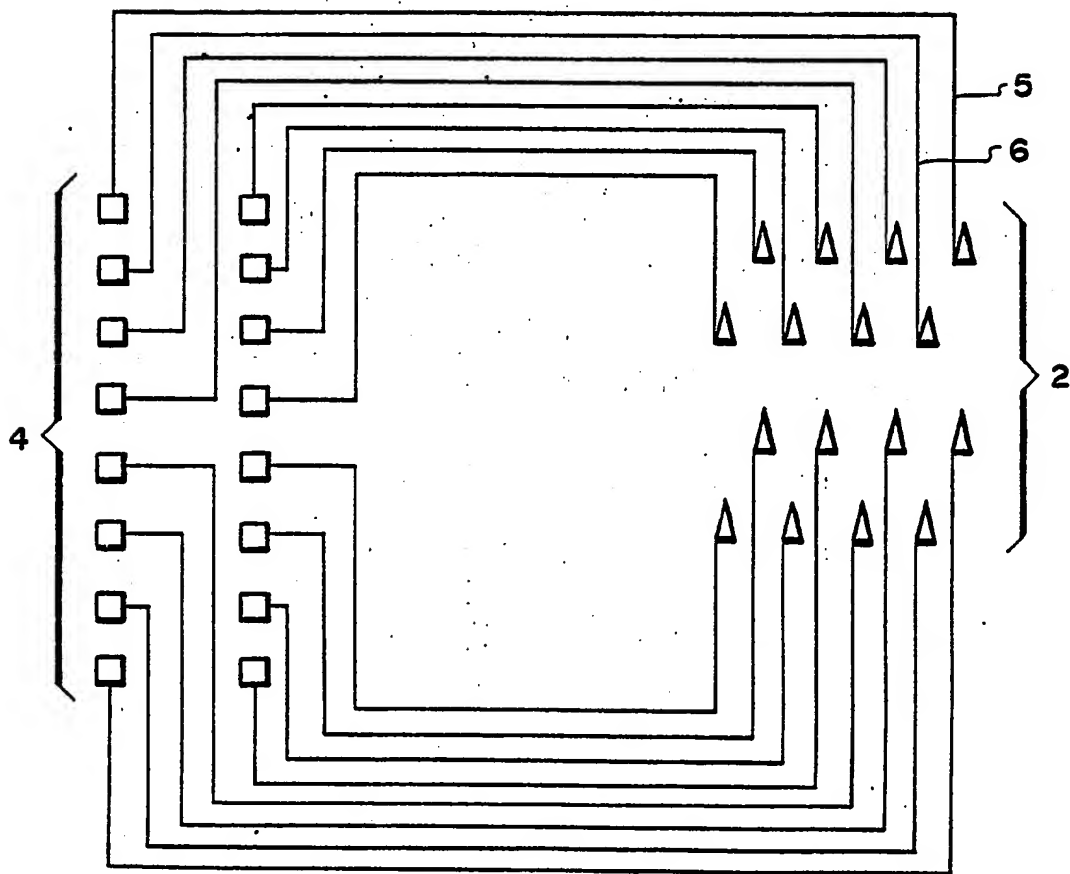


Fig. 2.

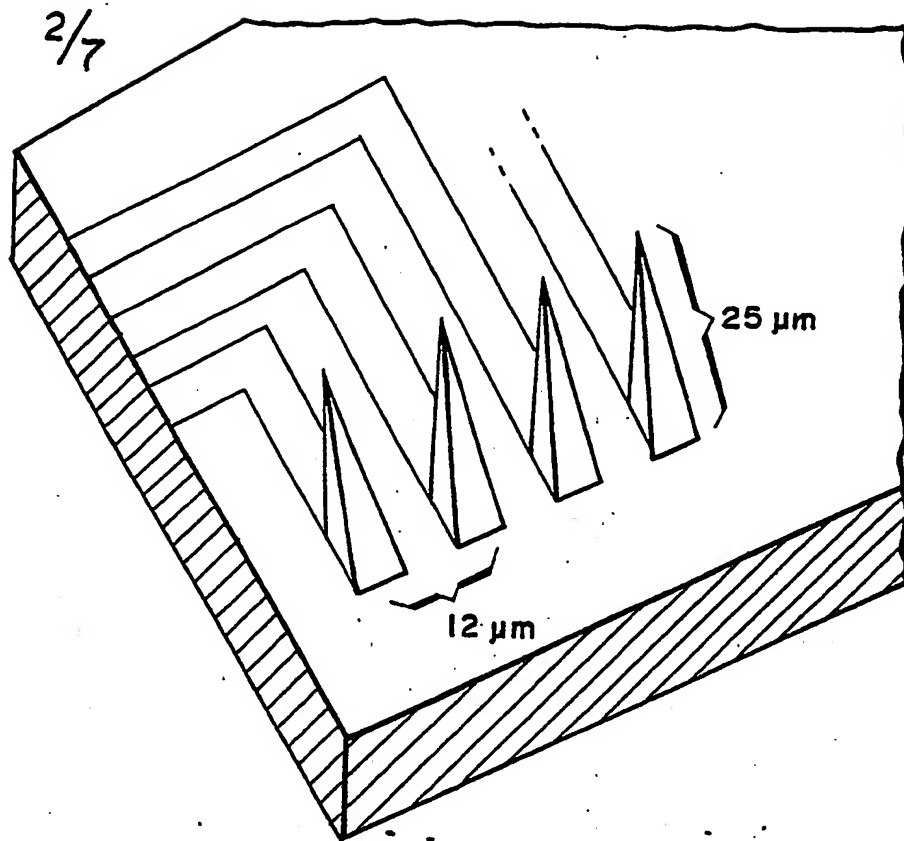


Fig. 3.

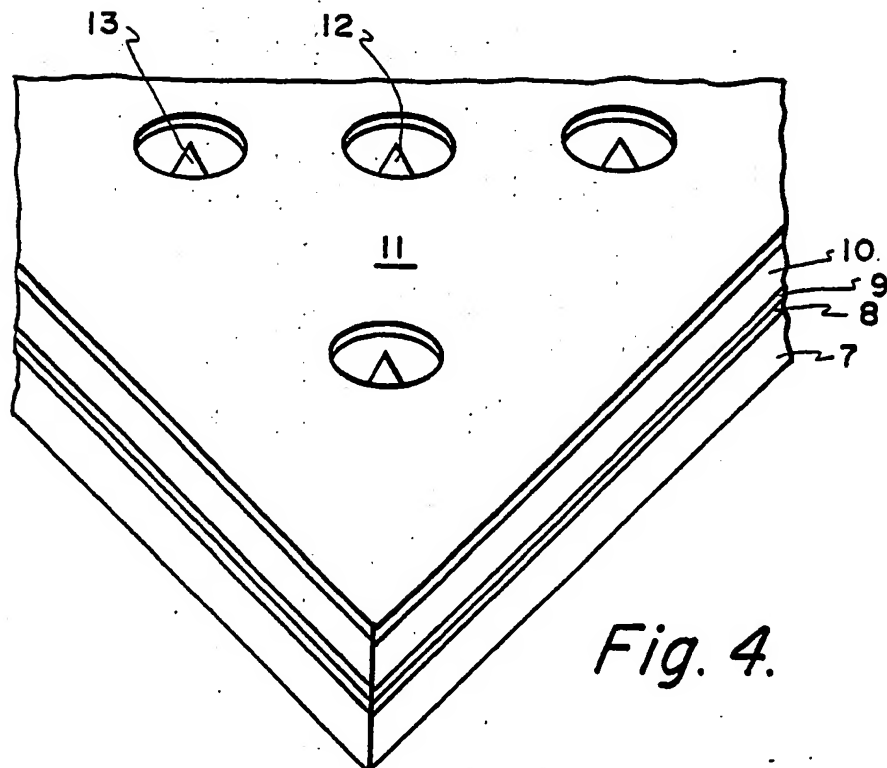


Fig. 4.

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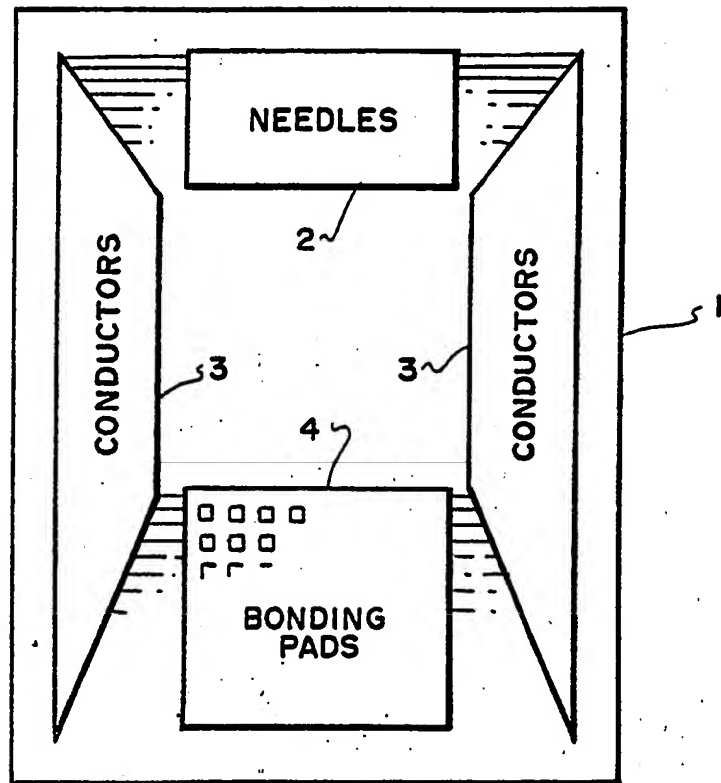


Fig. 5.

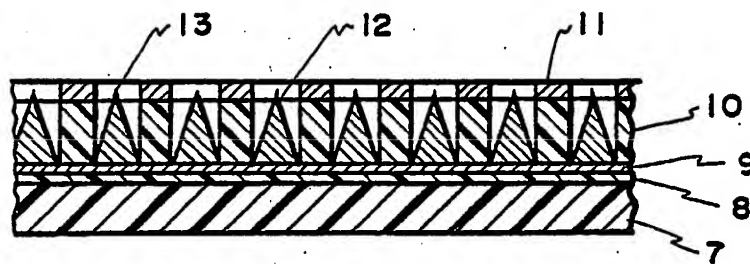


Fig. 6.

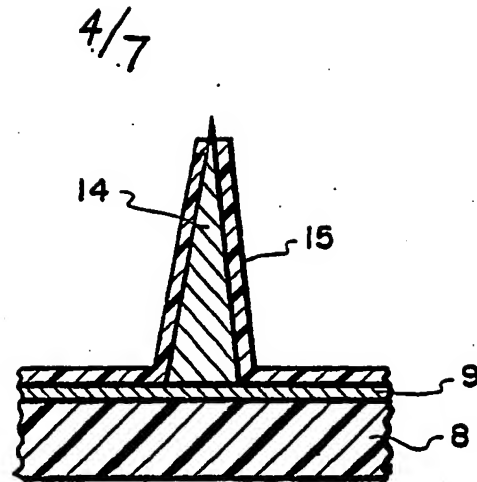


Fig. 7.

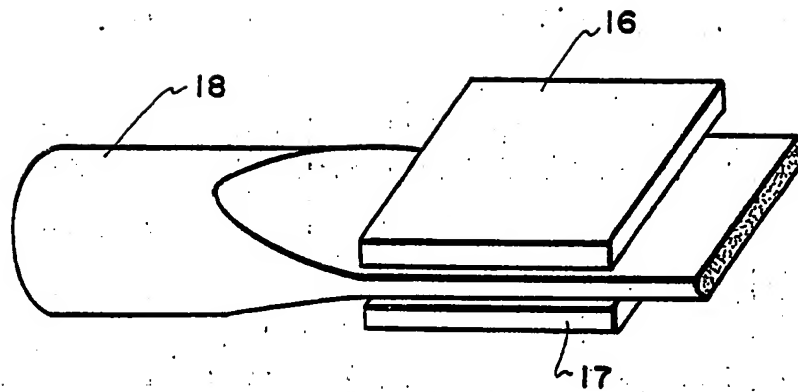


Fig. 8.

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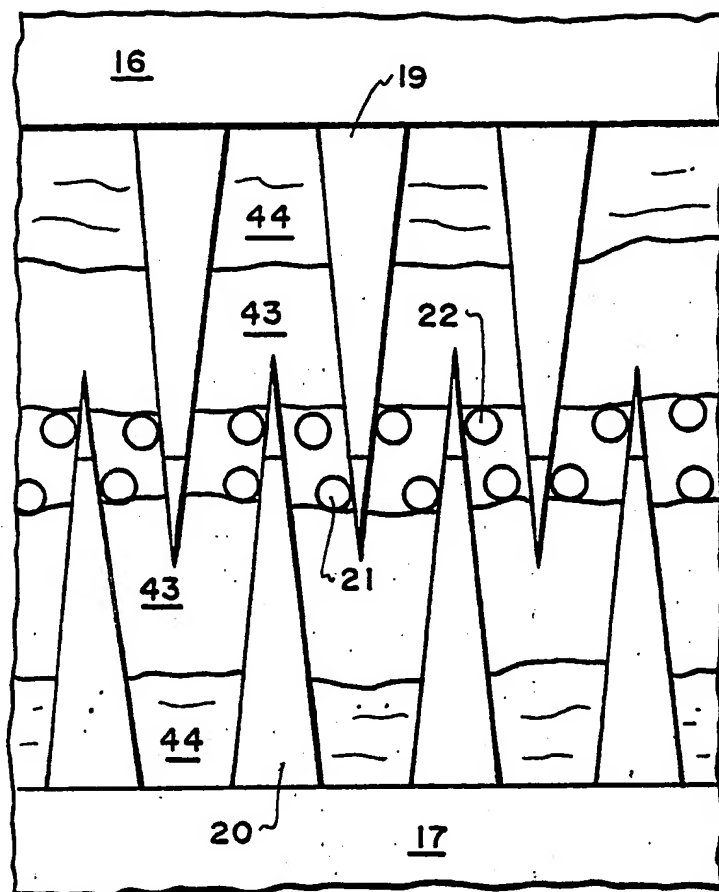


Fig. 9.

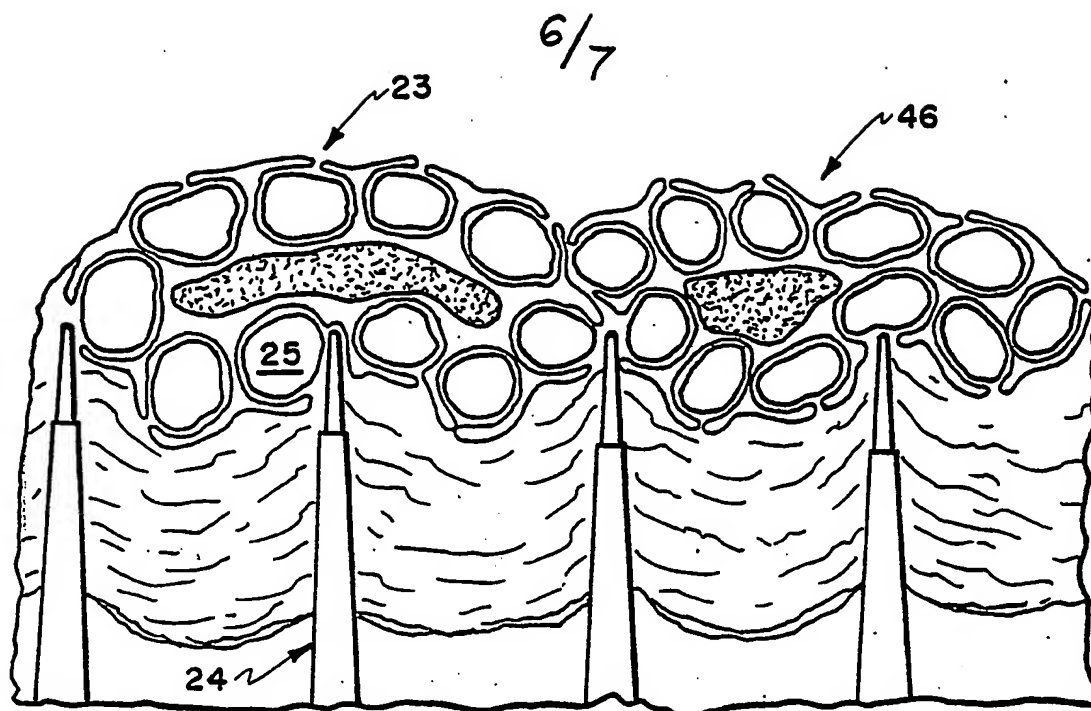


Fig. 10.

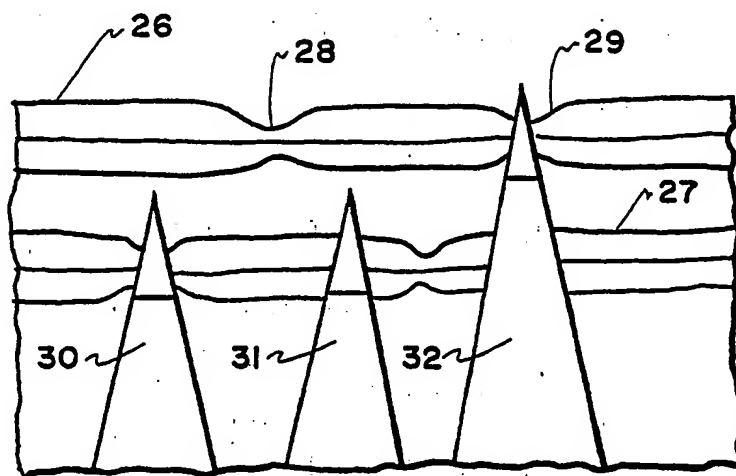


Fig. 11.

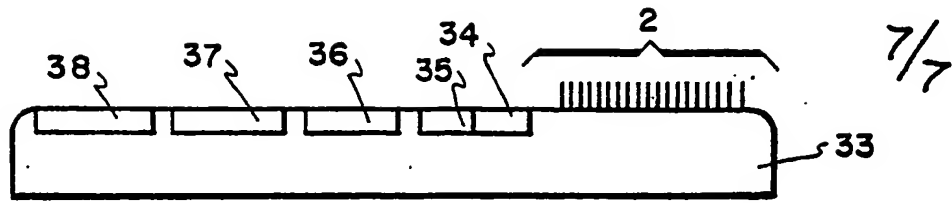


Fig. 12.

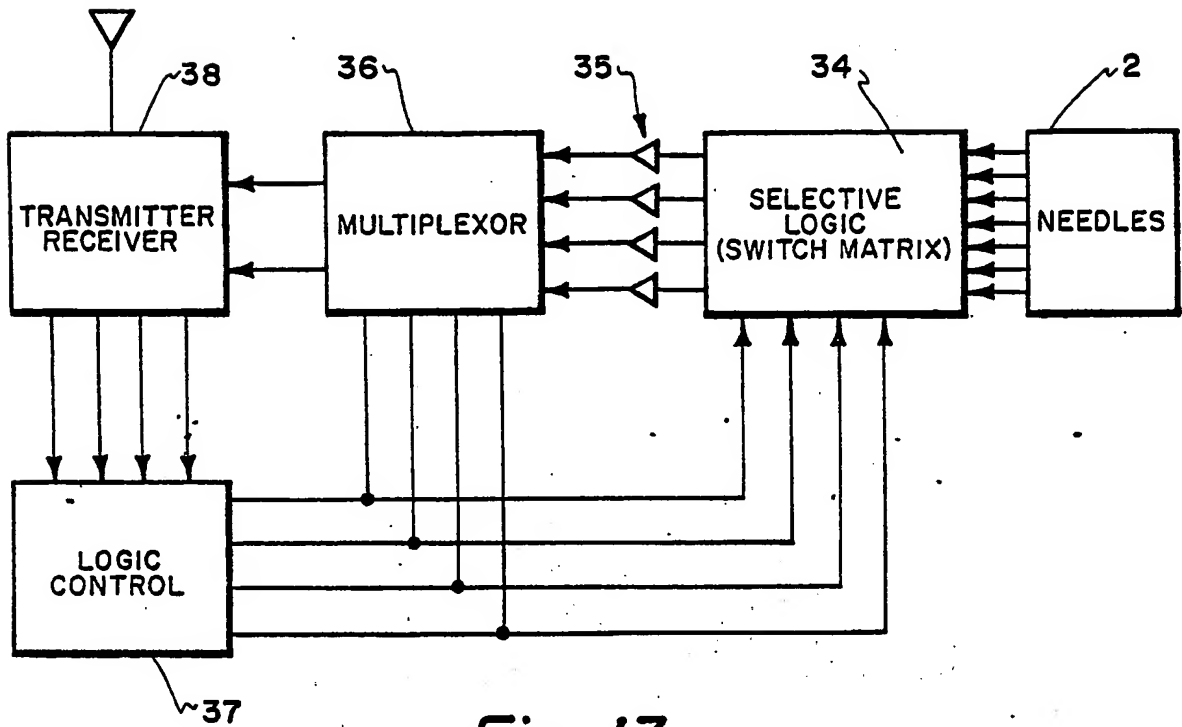


Fig. 13.

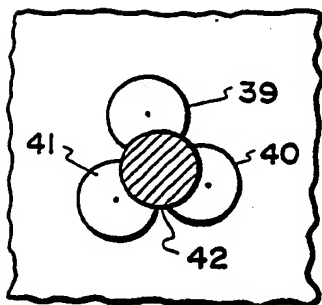


Fig. 14.

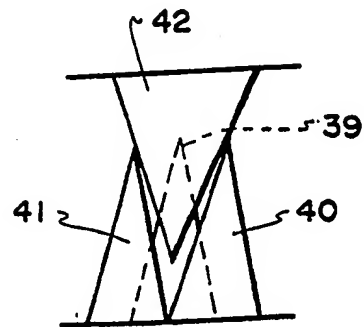


Fig. 15.

INTERNATIONAL SEARCH REPORT

International Application No

PCT/US87/01461

I. CLASSIFICATION OF SUBJECT MATTER (If several classification symbols apply, indicate all) *		
According to International Patent Classification (IPC) or to both National Classification and IPC		
IPC (4): A61B 5/04; A61N 1/05		
U.S. Cl. 128/642; 128/784		
II. FIELDS SEARCHED		
Minimum Documentation Searched *		
Classification System	Classification Symbols	
U.S.	128/639-642, 644, 784, 802 29/825	
Documentation Searched other than Minimum Documentation to the extent that such Documents are included in the Fields Searched *		
III. DOCUMENTS CONSIDERED TO BE RELEVANT **		
Category *	Citation of Document, ¹⁶ with indication, where appropriate, of the relevant passages ¹⁷	Relevant to Claim No. ¹⁸
X Y	IEEE Transactions On Bio-Medical Engineering, Volume BME-17, No. 3, issued July 1970, K. Wise, "An Integrated Circuit Approach to Extracellular Microelectrodes," see pages 238-246.	1-9, 19, 20, 34 10-16
X	US, A, 4,513,308 (GREENE) 23 April 1985 See the entire document.	29, 32
Y	Medical and Biological Engineering, Volume 12, No. 5, issued September 1974, D. Guyton, "Theory and design of capacitor electrodes for chronic stimulation," see pages 613-620.	10-16, 21, 22
A	DE, A, 2,558,281 (DEHNERT) 16 June 1977 See the entire document.	1-34
<p>* Special categories of cited documents: ¹⁵</p> <p>"A" document defining the general state of the art which is not considered to be of particular relevance</p> <p>"E" earlier document but published on or after the international filing date</p> <p>"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)</p> <p>"O" document referring to an oral disclosure, use, exhibition or other means</p> <p>"P" document published prior to the international filing date but later than the priority date claimed</p> <p>"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention</p> <p>"X" document of particular relevance: the claimed invention cannot be considered novel or cannot be considered to involve an inventive step</p> <p>"Y" document of particular relevance: the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.</p> <p>"d" document member of the same patent family</p>		
IV. CERTIFICATION		
Date of the Actual Completion of the International Search *		Date of Mailing of this International Search Report *
11 August 1987		13 SEP 1987
International Searching Authority *		Signature of Authorized Officer ¹⁹
ISA/US		Lee S. Cohen